

M.H.

13/4
152A

PCT/ZA 99/00094

P 023

Sertifikaat

REPUBLIEK VAN SUID-AFRIKA

Certificate

PATENTKANTOOR



REC'D 31 JAN 2000

PATENT OFFICE

DEPARTEMENT VAN HANDEL
EN NYWERHEID

REPUBLIC OF SOUTH AFRICA

DEPARTMENT OF TRADE
AND INDUSTRY

Hiermee word gesertifiseer dat
This is to certify that

ZA 99/94 4

The attached documents are true copies of the form P1, P6 and provisional specification of South African patent application No 98/9037

In the name of : **SASOL TECHNOLOGY (PROPRIETARY) LIMITED**
Filed on the : **05 OCTOBER 1998**
Entitled : **BIODEGRADABLE MIDDLE DISTILLATES AND
PRODUCTION THEREOF**

**PRIORITY
DOCUMENT**

SUBMITTED OR TRANSMITTED IN
COMPLIANCE WITH RULE 17.1(a) OR (b)

Geteken
Sign

PRETORIA

in die Republiek van Suid-Afrika, hierdie
in the Republic of South Africa, this

17th dag van SEPTEMBER 1999
day of

Registrateur van Patente
Registrar of Patents

REPUBLIC OF SOUTH AFRICA
PATENTS ACT, 1978
APPLICATION FOR A PATENT AND ACKNOWLEDGEMENT OF RECEIPT
[Section 30 (1) - Regulation 22]

Revenue stamps or revenue franking machine impression
Official date stamp

The grant of a patent is hereby requested by the undermentioned applicant on the basis of the present application filed in duplicate

OFFICIAL APPLICATION NO.		
21	01	989037

(i) APPLICANT'S OR AGENT'S REFERENCE
F190

FULL NAME(S) OF APPLICANT(S)

(ii)	71	SASOL TECHNOLOGY (PROPRIETARY) LIMITED a South African company
------	----	--

ADDRESS(ES) OF APPLICANT(S)

(iii)	1 STURDEE AVENUE, ROSEBANK, JOHANNESBURG REPUBLIC OF SOUTH AFRICA
-------	---

TITLE OF INVENTION

(iv)	54	BIODEGRADABLE MIDDLE DISTILLATES AND PRODUCTION THEREOF
------	----	--

(v)	The applicant claims priority as set out on the accompanying form P2. The earliest priority claimed is
-----	---

(vi)	This application is for a patent of addition to Patent Application No.	21	01	
------	--	----	----	--

(vii)	This application is a fresh application in terms of section 37 and is based on Patent Application No.	21	01	
-------	---	----	----	--

(viii)	This application is accompanied by:	<table border="0" style="width: 100%;"> <tr> <td style="width: 5%; text-align: center;"><input checked="" type="checkbox"/></td> <td style="width: 5%; text-align: center;">1.</td> <td style="padding: 2px;">A copy of a provisional or two copies of a complete specification of Provisional 23 pages.</td> </tr> <tr> <td style="text-align: center;"><input checked="" type="checkbox"/></td> <td style="text-align: center;">2.</td> <td style="padding: 2px;">Drawings of 1 sheets.</td> </tr> <tr> <td style="text-align: center;"><input checked="" type="checkbox"/></td> <td style="text-align: center;">3.</td> <td style="padding: 2px;">Publication particulars and abstract (form P8 in duplicate).</td> </tr> <tr> <td style="text-align: center;"><input checked="" type="checkbox"/></td> <td style="text-align: center;">4.</td> <td style="padding: 2px;">A copy of Figure 1 of the drawings for the abstract.</td> </tr> <tr> <td style="text-align: center;"><input type="checkbox"/></td> <td style="text-align: center;">5.</td> <td style="padding: 2px;">An assignment of invention.</td> </tr> <tr> <td style="text-align: center;"><input type="checkbox"/></td> <td style="text-align: center;">6.</td> <td style="padding: 2px;">Certified priority document(s) (state number):</td> </tr> <tr> <td style="text-align: center;"><input type="checkbox"/></td> <td style="text-align: center;">7.</td> <td style="padding: 2px;">Translation of the priority document(s).</td> </tr> <tr> <td style="text-align: center;"><input type="checkbox"/></td> <td style="text-align: center;">8.</td> <td style="padding: 2px;">An assignment of priority rights.</td> </tr> <tr> <td style="text-align: center;"><input type="checkbox"/></td> <td style="text-align: center;">9.</td> <td style="padding: 2px;">A copy of the form P2 and the specification of SA Patent Application No.</td> </tr> <tr> <td style="text-align: center;"><input checked="" type="checkbox"/></td> <td style="text-align: center;">10.</td> <td style="padding: 2px;">A declaration and power of attorney form P3.</td> </tr> </table>	<input checked="" type="checkbox"/>	1.	A copy of a provisional or two copies of a complete specification of Provisional 23 pages.	<input checked="" type="checkbox"/>	2.	Drawings of 1 sheets.	<input checked="" type="checkbox"/>	3.	Publication particulars and abstract (form P8 in duplicate).	<input checked="" type="checkbox"/>	4.	A copy of Figure 1 of the drawings for the abstract.	<input type="checkbox"/>	5.	An assignment of invention.	<input type="checkbox"/>	6.	Certified priority document(s) (state number):	<input type="checkbox"/>	7.	Translation of the priority document(s).	<input type="checkbox"/>	8.	An assignment of priority rights.	<input type="checkbox"/>	9.	A copy of the form P2 and the specification of SA Patent Application No.	<input checked="" type="checkbox"/>	10.	A declaration and power of attorney form P3.	21	01	
<input checked="" type="checkbox"/>	1.	A copy of a provisional or two copies of a complete specification of Provisional 23 pages.																																	
<input checked="" type="checkbox"/>	2.	Drawings of 1 sheets.																																	
<input checked="" type="checkbox"/>	3.	Publication particulars and abstract (form P8 in duplicate).																																	
<input checked="" type="checkbox"/>	4.	A copy of Figure 1 of the drawings for the abstract.																																	
<input type="checkbox"/>	5.	An assignment of invention.																																	
<input type="checkbox"/>	6.	Certified priority document(s) (state number):																																	
<input type="checkbox"/>	7.	Translation of the priority document(s).																																	
<input type="checkbox"/>	8.	An assignment of priority rights.																																	
<input type="checkbox"/>	9.	A copy of the form P2 and the specification of SA Patent Application No.																																	
<input checked="" type="checkbox"/>	10.	A declaration and power of attorney form P3.																																	

<input checked="" type="checkbox"/>	13.	Form P2 + copy
-------------------------------------	-----	----------------

(ix)	74	Address for service: HAHN & HAHN, 222 Richard Street, HATFIELD 0083, Pretoria
------	----	--

Dated this 05 day of October 19 98

Signature of applicant(s) or agent

This duplicate will be returned to the applicant's address for service as proof of lodging but is not valid unless endorsed with an official stamp

Received 1998-10-05 Official date stamp
REGISTRAR VAN PATENTE, MODELLE, HANDELSMERKE EN OUTEURSREG Registrar of Patents

REPUBLIC OF SOUTH AFRICA
PATENTS ACT, 1978
COMPLETE SPECIFICATION
[Section 30 (1) - Regulation 28]

OFFICIAL APPLICATION NO.		
21	01	989037

LODGING DATE	
22	05/10/98

INTERNATIONAL CLASSIFICATION	
51	C10L; B01J

FULL NAME(S) OF APPLICANT(S)

71

SASOL TECHNOLOGY (PROPRIETARY) LIMITED
a South African company

FULL NAME(S) OF INVENTOR(S)

72

ROBERT DE HAAN, LUIS PAPLO DANCUART,
MARK JAN PRINS, EWALD WATERMEYER DE WET

TITLE OF INVENTION

54

BIODEGRADABLE MIDDLE DISTILLATES AND PRODUCTION THEREOF

Biodegradable Middle Distillates and Production Thereof

5 This invention relates to middle distillates having biodegradability properties and to a process for production of such distillates. More particularly, this invention relates to middle distillates produced from a mainly paraffinic synthetic crude which is produced by the reaction of CO and H₂, typically by the Fischer-Tropsch (FT) process.

10 1. Background to the invention

In recent years a trend has developed to produce products which are so called "environmentally friendly", one aspect of which is biodegradability. To this end various bodies, such as ISO and the OECD have developed test methods to quantify
15 biodegradability. One such test is the CO₂ evolution test method, also known as the modified Sturm OECD method 301B, which test for ready biodegradability. In terms of this test, compounds can be considered to be readily biodegradable if they reach 60% biodegradation within 28 days.

20 Currently available middle distillates, typically crude oil derived diesel fuels, such as US 2-D grade (low sulphur No. 2-D grade for diesel fuel oil as specified in ASTM D 975-94) and/or CARB (California Air Resources Board 1993 specification) grade diesel, do not meet the biodegradability requirements of the abovementioned biodegradability test.

25

The prior art teaches in ZA 96/9890 that high biodegradability of hydrocarbon base oils could be derived from the presence of predominantly mono-methyl branching on the paraffinic carbon backbone. US 5,498,596 discloses a non-toxic, biodegradable

30 aromatics as well as other olefinic components. The biodegradability of the well fluid in the US patent can not be related back to the nature of the paraffinic molecules due to the fact that biodegradability is enhanced through branching and not through linear n-paraffinic molecules.

A need thus exists for a middle distillate cut, typically a diesel fuel, which is readily biodegradable as determined by the abovementioned biodegradability test.

- 5 Surprisingly, it has now been found, that a low aromatics content contributes to ready biodegradability of middle distillates, such as diesel fuel.

2. Summary of the invention

10

Thus, according to a first aspect of this invention, there is provided a biodegradable middle distillate cut, such as a diesel fuel, having an aromatics content of less than 9%, as determined by the ASTM D 5186 test method.

- 15 The synthetic middle distillate cut may have less than 8.99% (vol) monocyclic aromatics content.

The synthetic middle distillate cut may have less than 0.01% (vol) polycyclic aromatics.

20

The synthetic middle distillate cut may have an isoparaffins to n-paraffins mass ratio of between about 1:1 to about 12:1, typically the isoparaffins to n-paraffins mass ratio is between about 2:1 to about 6:1, and in one embodiment is 4:1.

-
- 25 The synthetic middle distillate cut may be a FT process product, or be at least partially produced in accordance with the FT process and/or process philosophy.

According to a second aspect of the invention, the synthetic middle distillate cut includes more than 50% isoparaffins, wherein the isoparaffins consist predominantly of methyl and/or ethyl and/or propyl branched isoparaffins.

- 5 At least some of the isoparaffins of the middle distillate cut may be methyl branched.

Typically, wherein at least some of the isoparaffins are di-methyl branched.

- 10 In a useful embodiment, at least 30% (mass) of the isoparaffins are mono-methyl branched.

Some of the isoparaffins may be ethyl branched, or even propyl branched.

- 15 According to a third aspect of the invention, there is provided a biodegradable synthetic middle distillate cut, having an aromatics content substantially as described above.

- 20 According to a fourth aspect of the invention, there is provided a biodegradable synthetic middle distillate cut, having an isoparaffinic content substantially as described above.

The invention extends to a biodegradable synthetic middle distillate cut having an isoparaffinic content and an aromatics content substantially as described above.

- 25 The biodegradable synthetic distillate may be a FT product.

According to a fifth aspect of the invention, there is provided a biodegradable diesel fuel composition including from 10% to 100% of a middle distillate cut as described above.

5

The biodegradable diesel fuel composition may include from 0 to 90% of another diesel fuel, such as conventional commercially available diesel fuel.

The biodegradable diesel fuel composition may include from 0 to 10% additives.

10

The additives may include a lubricity improver.

The lubricity improver may comprise from 0 to 0.5% of the composition, typically from 0.00001% to 0.05% of the composition. In a particularly useful embodiment, the

15

lubricity improver comprises from 0.008% to 0.02% of the composition.

The biodegradable diesel fuel composition may include a crude oil derived diesel, such as US 2-D grade diesel fuel and/or CARB grade diesel fuel, as the other diesel fuel of the composition.

20

According to yet another aspect of the invention, there is provided a process for producing a readily biodegradable synthetic middle distillate, the process including:

- (a) separating the products obtained from synthesis gas via the FT synthesis reaction into one or more heavier fraction and one or more lighter fraction;
- 25- (b) catalytically processing the heavier fraction under conditions which yield mainly middle distillates;

- (c) separating the middle distillate product of step (b) from a light product fraction and a heavier product fraction which are also produced in step (b); and
- (d) blending the middle distillate fraction obtained in step (c) with at least a portion of the one or more lighter fraction of step (a), or products thereof.

5

The catalytic processing of step (b) may be a hydroprocessing step, for example, hydrocracking.

10

The process for producing a synthetic middle distillate may include one or more additional step of fractionating at least some of the one or more lighter fraction of step (a), or products thereof, prior to step (d).

15

The process for producing a synthetic middle distillate may include the additional step of hydrotreating at least some of the one or more light fraction of step (a), or products thereof, prior to step (d).

The one or more heavier fraction of step (a) may have a boiling point above about 270°C, however, it may be above 300°C.

20

The one or more lighter fraction may have a boiling point in the range C₅ to the boiling point of the heavier fraction, typically in the range 160°C to 270°C.

25

The product of step (d) may boil in the range 100°C to 400°C. The product of step (d) may boil in the range 160°C to 370°C.

The product of step (d) may be obtained by mixing the middle distillate fraction obtained in step (c) with at least a portion of the one or more lighter fraction of step (a), or products thereof, in a volume ratio of between 1:1 and 9:1, typically 2:1 and 6:1, and in one embodiment, in a volume ratio of 84:16.

The product of the above process may be a synthetic middle distillate cut, or products thereof, or compositions thereof, as described above.

The product of step (d) may be a diesel fuel.

A biodegradable diesel fuel produced in accordance with this invention may be produced from a mainly paraffinic synthetic crude (syncrude) obtained from synthesis gas (syngas) through a reaction like the FT reaction.

5

The FT products cover a broad range of hydrocarbons from methane to species with molecular masses above 1400; including mainly paraffinic hydrocarbons and much smaller quantities of other species such as olefins and oxygenates. Such a diesel fuel could be used on its own or in blends to improve the quality of other diesel fuels not meeting the current and/or proposed, more stringent fuel quality and environmental specifications.

10

The invention extends to an essentially non-polluting, readily biodegradable diesel fuel composition comprising of a mixture of normal paraffins (n-paraffins) and iso-paraffins in the typical diesel range from 160-370°C, having an iso-paraffin:n-paraffin mass ratio from about 2:1 to about 12:1, more typically from 2:1 to 6:1, and the iso-paraffins of the mixture contain greater than 30%, based on the total mass of the iso-paraffins in the mixture, of mono-methyl species, with the balance consisting mainly of ethyl and/or dimethyl branched species. These iso-paraffins contained in a mixture with minor amounts of aromatics and other materials, contribute to a product from which readily biodegradable diesel fuels can be obtained.

15

20

This diesel will readily biodegrade in an aquatic environment under aerobic conditions. This biodegradability can be attributed to the very low aromatic content present in the middle distillate cut, typically a diesel fuel. The aromatic content will typically comprise 2.5% (mass) of monocyclic, 0.2% (mass) of dicyclic and <10 ppm (mass) of polycyclic aromatics with a total aromatic content of around 2.7% (mass).

25

3. Specific Description

Process

- 5 The process of this invention provides a process for the conversion of primary FT products into naphtha and middle distillates, specifically high performance diesel.

10 The FT process is used industrially to convert synthesis gas, derived from coal, natural gas, biomass or heavy oil streams, into hydrocarbons ranging from methane to species with molecular masses above 1400. While the main products are linear paraffinic materials, other species such as branched paraffins, olefins and oxygenated components form part of the product slate. The exact product slate depends on reactor configuration, operating conditions and type of catalyst that is employed, as is evident from e.g. Catal.Rev.-Sci. Eng., 23(1&2), 265-278 (1981).

15 Typical reactors for the production of heavier hydrocarbons (i.e. waxy hydrocarbons) are the Slurry Bed or the Tubular Fixed Bed types, while typical operating conditions are 160 – 280°C, in some cases 210-260°C, and 18 – 50 Bar, in some cases 20-30 Bar. Active metals typically useable in the catalyst used in such a reactor include
20 iron, ruthenium or cobalt. While each catalyst will give its own unique product slate, in all cases the product contains some waxy, highly paraffinic material which needs to be further upgraded into usable products. The FT products' can be converted into a range of final products, such as middle distillates, gasoline, solvents, lube oil bases, etc. Such conversion, which usually consists of a range of processes such
25 as hydrocracking, hydrotreatment and distillation, can be termed a FT work-up process.

~~The FT work-up process of this invention uses a feed stream consisting of C₅ and higher hydrocarbons derived from a FT process. This feed is separated into at least two fractions, a heavier and at least one lighter fraction. The cut point between the two fractions is usually less than 300°C and typically around 270°C.~~

The table below gives a typical composition of the two fractions, within about 10% accuracy:

Table 1 - Typical Fischer-Tropsch product after separation into two fractions

Boiling range	Condensate (< 270°C, volume %)	Wax (> 270°C, volume %)
C ₅ -160°C	45	
160-270°C	51	3
270-370°C	4	35
370-500°C		42
> 500°C		20

The >270°C fraction, also referred to as wax, contains a considerable amount of hydrocarbon material, which boils higher than the normal diesel range. If we consider a diesel boiling range of 100-400°C, typically 160-370 °C, it means that all material heavier than about 370°C needs to be converted into lighter materials by means of a catalytic process often referred to as hydrocracking. Catalysts for this step are of the bifunctional type; i.e. they contain sites active for cracking and for hydrogenation. Catalytic metals active for hydrogenation include group VIII noble metals, such as platinum or palladium, or sulphided Group VIII base metals, e.g. nickel, cobalt, which may or may not include a sulphided Group VI metal, e.g. molybdenum. The support for the metals can be any refractory oxide, such as silica, alumina, titania, zirconia, vanadia and other Group III, IV, VA and VI oxides, alone or in combination with other refractory oxides. Alternatively, the support can partly or totally consist of zeolite. Amorphous silica-alumina is the preferred support for middle distillates conversion.

Process conditions for hydrocracking can be varied over a wide range and are usually laboriously chosen after extensive experimentation to optimise the yield of middle distillates. In this regard, it is important to note that, as in many chemical reactions, there is a trade-off between conversion and selectivity. A very high conversion will result in a high yield of gases and low yield of distillate fuels. It is therefore important to painstakingly tune the process conditions in order to limit the conversion of >370°C hydrocarbons. Table 2 lists some of the conditions found, after extensive experimentation, to provide a desirable product range.

Table 2: - Typical Hydrocracking Process Conditions

Process Condition	Broad Range	Preferred Range
Temperature, °C	150-450	340-400
Pressure, bar(g)	10-200	30-80
Hydrogen Flow Rate, m ³ /m ³ feed	100-2000	800-1600
Conversion of >370°C material, Mass %	30-80	50-70

5 It will be clear to those skilled in the art that it is possible to convert all the >370°C material in the feedstock by recycling the part that is not converted during the hydrocracking process.

10 As is evident from table 1, most of the fraction boiling below 270°C is already boiling in the typical boiling range for diesel, i.e. 160-370°C. This fraction may or may not be subjected to hydrotreating. By hydrotreating, heteroatoms are removed and unsaturated compounds are hydrogenated. Hydrotreating is a well-known industrial process catalysed by any catalyst having a hydrogenation function, e.g. Group VIII noble metal or a sulphided base metal or sulphided Group VI metals, or combinations thereof. Preferred supports are alumina and silica. Table 3 lists typical operating conditions for the hydrotreating process.

15

Table 3 - Typical Hydrotreating Process Conditions

Process Condition	Broad Range	Preferred Range
Temperature, °C	150-450	200-400
Pressure, bar(g)	10-200	30-80
Hydrogen Flow Rate, m ³ /m ³ feed		

While the hydrotreated fraction may be fractionated into paraffinic materials useful as solvents, the applicant has now found that the hydrotreated fraction may be directly blended with the products obtained from hydrocracking the wax. Although it is possible to hydroisomerise the material contained in the condensate stream, the

5 applicant has found that this leads to a small, but significant loss of material in the diesel boiling range to lighter material. Furthermore, isomerisation leads to the formation of branched isomers, which leads to Cetane ratings less than that of the corresponding normal paraffins (n-paraffins).

10 Several diesel fuels, produced broadly in accordance with the invention, as well as other crude oil derived diesel fuels such as US 2-D grade and CARB grade, were tested by the applicant. The basic characteristics of the fuels tested for biodegradability are included in Table 4(a).

15 Synthetic diesel fuels, produced broadly in accordance with this invention, and other conventional diesels were tested by the applicant. It was found that there were significant differences regarding the chemical composition of the fuels.

In particular, the synthetic fuels contained very small quantities of aromatic species.

20 Other differences relate to the predominance of paraffinic species in the synthetic diesels, as can be seen from Table 4(b).

Upon analysis, it thus appears, since most of the other characteristics of the synthetic and conventional diesel fuels are not very dissimilar, the difference in the

25 biodegradability performance can be attributed to the differences in the chemical nature indicated above.

Table 4(a) - Basic Characteristics of the Tested Fuels

Fuel Name		SPD Diesel Type A	SPD Diesel Type B	Commercial US 2D	CARB* Protocol Standard
Fuel Code		S1	S2	P1	P2
Density (15°C)	Kg/dm ³	0.7769	0.7779	0.8547	0.8308
Distillation	ASTM D86				
IBP	°C	189	185	184	203
10%	°C	209	208	214	218
50%	°C	256	257	259	249
90%	°C	331	332	312	290
FBP	°C	356	358	342	351
SFC Aromatics (mass %)	ASTM D5186	2.68%	2.46%	38.22%	9.91%
Monocyclic		93.61%	N/A	71.35%	99.54%
Bicyclic		6.38%	N/A	26.54%	0.45%
Polycyclic		0.01%	N/A	2.11%	0.01%
Sulphur (mass %)	ASTM D4294	0.001%	0.002%	0.022%	0.028%

5 * CARB – California Air Resources Board

Furthermore, in a specific middle distillate produced in accordance with this invention, the total amount of isoparaffins in the light boiling range of the diesel (160-

10 following Table 4(b).

Table 4(b) – Isoparaffins:n-Paraffins of Middle Distillate Fractions

Boiling Range	Corresponding Carbon Range	Average Iso:Normal Paraffins Ratio	
		Range	Typical value
160-270°C	C ₁₀ -C ₁₇	0.5 - 4.0	2.2
270-370°C	C ₁₇ -C ₂₃	4.0 - 14.0	10.5

- 5 It is this unique composition of the synthetic fuel, which is directly caused by the way in which the FT work-up process of this invention is operated, that contributes to the unique characteristics of said middle distillates.

10 The applicant has also found, that from the perspective of fuel quality, it is not necessary to hydrotreat the <270°C fraction, adding said fraction directly to the products from hydrocracking the wax. While this results in the inclusion of oxygenates and unsaturates in the final diesel, fuel specifications usually allow for this. Circumventing the need for hydrotreatment of the condensate results in considerable savings of both capital and operating cost.

15 The invention will now be illustrated, by way of non-limiting examples only, with reference to the accompanying Figure 1.

20 A FT work-up process is outlined in the attached Figure 1. The synthesis gas (syngas), a mixture of Hydrogen and Carbon Monoxide, enters the FT reactor 1 where the synthesis gas is converted to hydrocarbons by the FT process.

A lighter FT fraction is recovered in line 7, and may or may not pass through fractionator 2 and hydrotreater 3. The product 9 (9a) from the hydrotreater may be
 25 separated in fractionator 4 or, alternatively, mixed with hydrocracker 5 products 16 and sent to a common fractionator 6.

A waxy FT fraction is recovered in line 13 and sent to hydrocracker 5. If fractionation 2 is considered then the bottoms cut 12 are also sent to hydrocracker 5. The products 16, on their own or mixed with the lighter fraction 9a, are separated in fractionator 6.

5

Depending on the process scheme, a light product fraction, naphtha 19, is obtained from fractionator 6 or by blending equivalent fractions 10 and 17. This is a C₅-160°C fraction useful as naphtha.

10 A somewhat heavier cut i.e. the middle cut, synthetic diesel 20, is obtainable in a similar way from fractionator 6 or by blending equivalent fractions 11 and 18. This cut is recovered as a 160-370°C fraction useful as diesel

The heavy unconverted material 21 from fractionator 6 is recycled to extinction to
15 hydrocracker 5. Alternatively, the residue may be used for production of synthetic lube oil bases. A small amount of C₁-C₄ gases is also separated in fractionator 6.

The described FT work-up process of Figure 1 may be combined in a number of configurations. The applicant considers these an exercise in what is known in the art
20 as Process Synthesis Optimisation.

However, the specific process conditions for the Work-up of Fischer-Tropsch primary products, the possible process configurations of which are outlined in Table 5, were obtained after extensive and laborious experimentation and design.

25

Table 5 - Possible Fischer-Tropsch Product Work-up Process Configurations

Process Step			Process Configuration					
			A	B	C	D	E	F
2	Light FT Product Fractionator				X			X
4	Hydrotreater	Products		X			X	X
	Fractionator							
5	Waxy FT	Product	X	X	X	X	X	X
	Hydrocracker							
6	Hydrocracked	Products	X	X	X	X	X	X
	Fractionator							

Number	Reference numerals of Figure 1
FT	Fischer-Tropsch

5 4. Experimental Procedure

The biodegradability of the fuels was tested using the Carbon Dioxide Evolution method (modified Sturm OECD Method 301B). This method tests for ready biodegradability. A compound can be considered readily biodegradable if it reaches
10 60% biodegradation within 28 days under the prescribed test conditions. Domestic activated sludge, not previously exposed to industrial effluent, was used as the source of micro-organisms for the test. The biodegradability tests were continuously validated using Sodium acetate as a reference chemical for checking the viability of the micro-organisms.

15

The test involves aerating the sample by passing carbon dioxide-free air at a controlled rate in the dark or in diffuse light. The sample must be the only source of carbon. Degradation is followed over 28 days by determining the carbon dioxide produced. This gas is trapped in barium or sodium hydroxide, and it is measured by
20 titration of the residual hydroxide or as inorganic carbon. For additional details refer to the standard procedure.

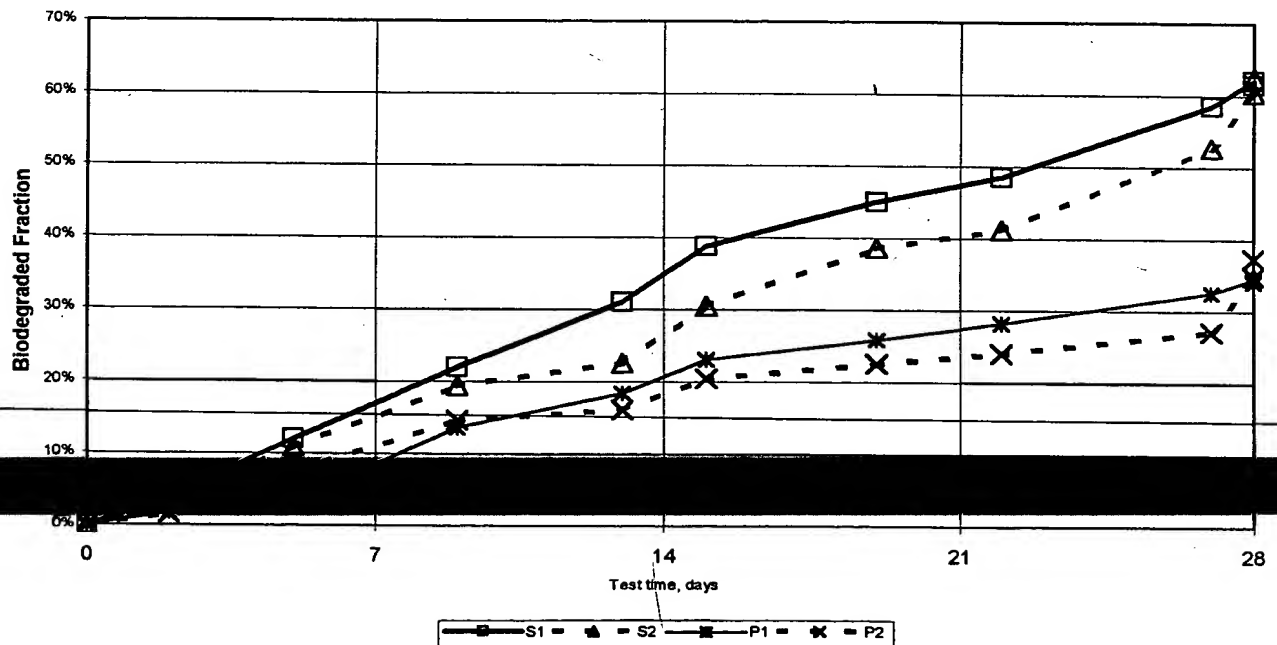
The results of the tests are set out in table 6 and chart 1 below.

Table 6: Biodegradability of Diesel Fuels (Modified Sturm Test)

Days from start of test sequence	Synthetic Diesels		Petroleum Diesels	
	SPD A	SPD B	US 2D	CARB
	S1	S2	P1	P2
0	0%	0%	0%	0%
2	4%	4%	2%	2%
5	12%	11%	4%	7%
9	22%	19%	14%	15%
13	31%	23%	18%	16%
15	39%	30%	23%	20%
19	45%	39%	26%	22%
22	48%	41%	28%	24%
27	58%	53%	32%	27%
28	62%	60%	34%	35%
28	61%	63%	34%	37%

NB. Two results are available at the end of the test sequence after acidification of the test media

Chart 1: Biodegradability Test Results (Modified Sturm Test)



5. Examples

Example 1

5 Fuel S1 was produced broadly in accordance with the invention, by following the process described above. It is a fully hydroprocessed fuel. The fractionation of the two basic components was completed in separate steps. S1 diesel was a blend of 84% (vol) of hydrocracked diesel (product stream 11 from fractionator 4) and 16% (vol) of hydrotreated diesel (product stream 18 from fractionator 6) produced using configuration B of Table 5. It contained 2.68% total aromatics, most of the aromatics species being monocyclic.

15 This fuel biodegraded 61% after 28 days under the conditions specified for the described modified Sturm OECD Method 301B. A fuel with this behaviour is considered biodegradable.

Example 2

20 Fuel S2 was produced by hydrocracking of the FT wax and distilling the diesel fraction (product stream 18). The primary light FT products were distilled separately (product stream 11 produced without passing through hydrotreater 3). S2 diesel was obtained by blending these two cuts in a 84:16 ratio (volume). Process Configuration C of Table 5 was used to produce this fuel. The total aromatics content was 2.46%.

25 This fuel biodegraded 63% after 28 days under the same conditions described in example 1. This fuel can also be considered biodegradable.

Example 3

30

Fuel P1 is a commercial diesel procured in the United States of America. It meets the US 2D diesel specification. This conventional petroleum based diesel fuel contained 38,22% aromatics, almost 71% of which were monocyclic species.

This fuel biodegraded 34% under the conditions described in example 1. A fuel with this behaviour is not considered biodegradable.

Example 4

5

Fuel P2 is a non-commercial fuel procured in the United States of America. It meets the specifications of the California Air Resources Board (CARB) protocol. This fuel contained 9,91% aromatics, mainly monocyclic species. In spite of this, this fuel biodegraded only ca 37% under the conditions described in example 1.

10

A fuel with this behaviour is not considered biodegradable.

Claims:

1. A synthetic middle distillate cut having less than 9 mass% , according to ASTM D 5186, aromatics content.
2. A synthetic middle distillate cut as claimed in claim 1, having less than 8.99 mass% monocyclic aromatics content.
3. A synthetic middle distillate cut as claimed in claim 1 or claim 2, having less than 0.01 mass% polycyclic aromatics.
4. A synthetic middle distillate cut as claimed in any one of the preceding claims, having an isoparaffins to n-paraffins mass ratio of between about 1:1 to about 12:1.
5. A synthetic middle distillate cut as claimed in claim 4, wherein the isoparaffins to n-paraffins mass ratio is between about 2:1 to about 6:1.
6. A synthetic middle distillate cut as claimed in claim 5, wherein the isoparaffins to n-paraffins mass ratio is 4:1.
7. A synthetic middle distillate cut as claimed in any one of the preceding claims, wherein the synthetic distillate is derived from a FT primary product.
8. A synthetic middle distillate cut comprising more than 50% isoparaffins, wherein the isoparaffins are predominantly methyl and/or ethyl and/or propyl branched.
9. A middle distillate cut as claimed in claim 8, wherein at least some of the isoparaffins are methyl branched.
10. A middle distillate cut as claimed in claim 8 or claim 9, wherein at least some of the isoparaffins are di-methyl branched.
11. A middle distillate cut as claimed in any one of claims 8 to 10, wherein at least 30% (mass) of the isoparaffins are mono-methyl branched.

12. A middle distillate cut as claimed in any one of claims 8 to 11, wherein at least some of the isoparaffins are ethyl branched.
13. A biodegradable synthetic middle distillate cut, having an aromatics content substantially as claimed in any one of claims 1 to 7.
- 5 14. A biodegradable synthetic middle distillate cut, having an isoparaffinic content substantially as claimed in any one of claims 8 to 12.
15. A biodegradable synthetic middle distillate cut, having an isoparaffinic content as claimed in claim 13 and an aromatics content as claimed in claim 14.
- 10 16. A synthetic middle distillate cut as claimed in any one of claims 8 to 15, wherein the synthetic distillate is a FT product.
17. A biodegradable diesel fuel composition including from 10% to 100% of a middle distillate cut as claimed in any one of the preceding claims.
18. A biodegradable diesel fuel composition as claimed in claim 17, including from 0 to 90% of at least one other diesel fuel.
- 15 19. A biodegradable diesel fuel composition as claimed in claim 17 or claim 18, including from 0 to 10% additives.
20. A biodegradable diesel fuel composition as claimed in any one of claims 17 to 19, wherein the additives include a lubricity improver.
- 20 21. A biodegradable diesel fuel composition as claimed in claim 20, wherein the lubricity improver comprises from 0 to 0.5% of the composition.
22. A biodegradable diesel fuel composition as claimed in claim 21, wherein the
-
23. A biodegradable diesel fuel composition as claimed in claim 22, wherein the lubricity improver comprises from 0.008% to 0.02% of the composition.

24. A biodegradable diesel fuel composition as claimed in any one of claims 18 to 23, wherein one of the other diesel fuels is US 2-D grade diesel fuel.

25. A biodegradable diesel fuel composition as claimed in any one of claims 18 to 23, wherein one of the other diesel fuels is CARB grade diesel fuel.

5 26. A process for producing a readily biodegradable synthetic middle distillate, the process including:

(a) separating the products obtained from synthesis gas via the FT synthesis reaction into one or more heavier fraction and one or more lighter fraction;

10 (b) catalytically processing the one or more heavier fraction under conditions which yield mainly middle distillates;

(c) separating the middle distillate product of step (b) from the lighter product and heavier product that are also produced in step (b); and

15 (d) blending the middle distillate fraction obtained in step (c) with at least a portion of the one or more lighter fraction of step (a), or products thereof.

27. A process for producing a synthetic middle distillate as claimed in claim 26, wherein the catalytic processing of step (b) is a hydroprocessing step.

20 28. A process for producing a synthetic middle distillate as claimed in claim 26 or claim 27, including one or more additional step of fractionating at least some of the one or more lighter fraction of step (a), or products thereof, prior to step (d).

25 29. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 28, including the additional step of hydrotreating at least some of the one or more light fraction of step (a), or products thereof, prior to step (d).

30. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 29, wherein the one or more heavier fraction of step (a) boils above about 270°C.

30- 31. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 30, wherein the one or more heavier fraction of step (a) boils above about 300°C.

32. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 31, wherein the one or more lighter fraction boils in the range C₅ to the boiling point of the heavier fraction.
- 5 33. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 32, wherein the one or more lighter fraction boils in the range 160°C to 270°C.
34. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 33, wherein the product of step (d) boils in the range 100°C to 400°C.
- 10 35. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 34, wherein the product of step (d) boils in the range 160°C to 370°C.
36. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 35, wherein the product of step (d) is a diesel fuel.
- 15 37. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 36, wherein the product of step (d) is readily biodegradable.
38. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 37, wherein the product of step (d) is obtained by mixing the middle distillate fraction obtained in step (c) with at least a portion of the one or more lighter fraction of step (a), or products thereof, in a volume ratio selected to provide a diesel fuel having a required specification.
- 20 39. A process for producing a synthetic middle distillate as claimed in any one of claims 26 to 38, wherein the product of step (d) is obtained by mixing the middle distillate fraction obtained in step (c) with at least a portion of the one or more lighter fraction of step (a), or products thereof, in a volume ratio of between 1:1 and 9:1.
- 25 40. A process for producing a synthetic middle distillate as claimed in claim 39, wherein the product of step (d) is obtained by mixing the middle distillate fraction obtained in step (c) with at least a portion of the one or more lighter fraction of step (a), or products thereof, in a volume ratio of between 2:1 and 6:1.

41. A process for producing a synthetic middle distillate as claimed in claim 40, wherein the product of step (d) is obtained by mixing the middle distillate fraction obtained in step (c) with at least a portion of the one or more lighter fraction of step (a), or products thereof, in a volume ratio of 84:16.

42. A synthetic middle distillate cut, substantially as herein described and illustrated.

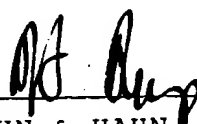
43. A biodegradable synthetic middle distillate cut, substantially as herein described and illustrated.

44. A biodegradable diesel fuel composition, substantially as herein described and illustrated.

45. A process for producing a readily biodegradable synthetic middle distillate, substantially as herein described and illustrated.

46. A new synthetic middle distillate cut, biodegradable synthetic middle distillate cut, biodegradable diesel fuel composition, or a new process for producing a readily biodegradable synthetic middle distillate, substantially as herein described.

DATED THIS 5th DAY OF OCTOBER 1998


HAHN & HAHN
Agent for Applicant

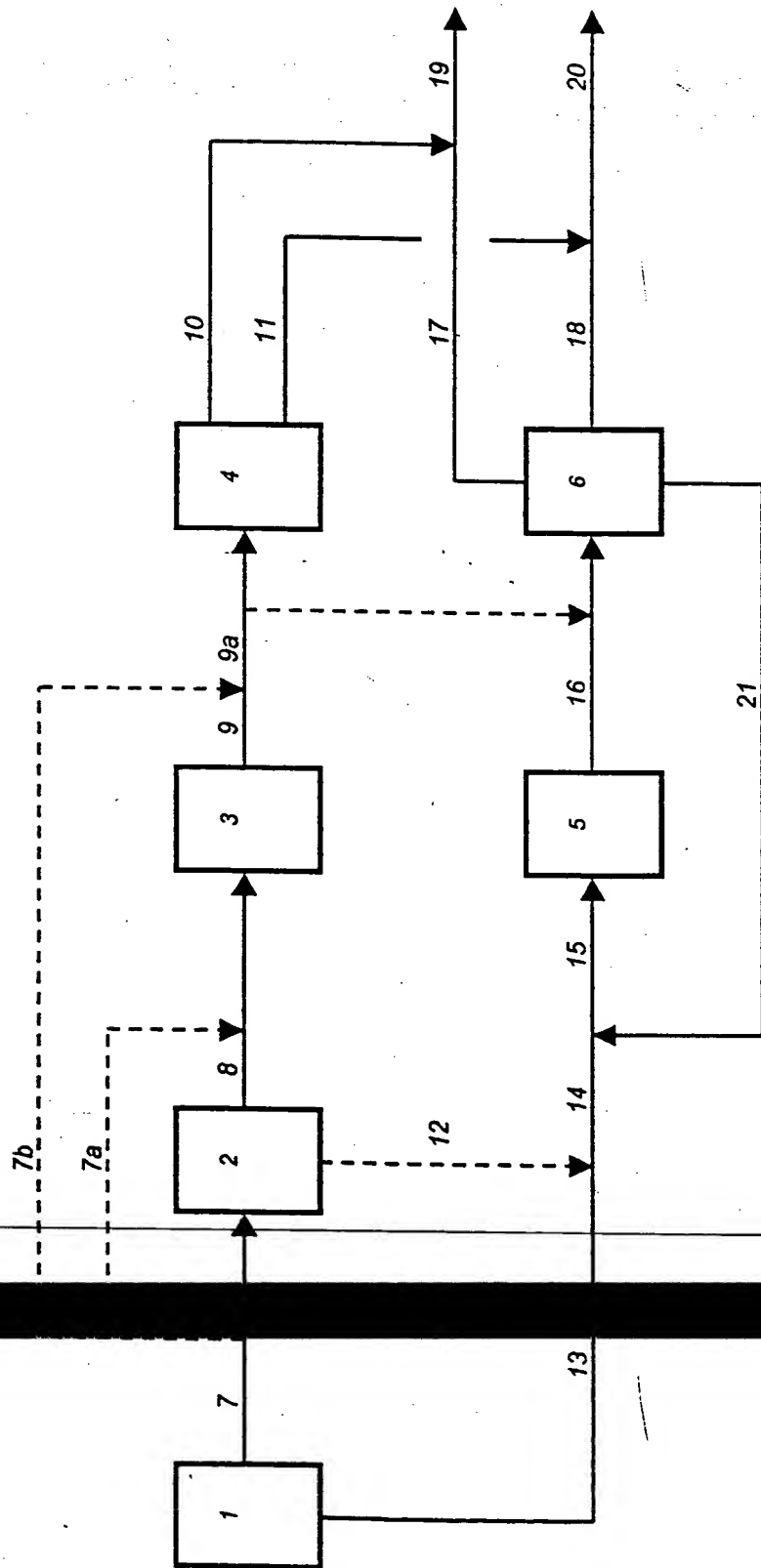


Figure 1